This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) Compounds of the formula I

$$R^{1}$$
 $R^{1/2}$
 $R^{1/2}$
 $R^{1/2}$
 $R^{1/2}$
 $R^{1/2}$

in which

X = N or CH,

 R^1 , R^2 , R^3 = independently of one another OH, OA, CN, Hal, COR^4 or CH_2R^4 , R^4 = OH, OA, NH₂, NHB or NB₂,

 $Q = CH_{27}$ or CO and, if divalent, also CH,

A, B = independently of one another straight-chain or branched alkyl or alkoxy having 1 to 10 C atoms, alkenyl having 2 to 10 C atoms or alkoxyalkyl having 2 to 10 C atoms,

m = 2, 3, 4, 5 or 6 and

n = 0, 1, 2, 3 or 4,

andor physiologically acceptable salts, derivatives, solvates andor stereoisomers thereof, including mixtures thereof in all ratios.

2. (Currently Amended) Compounds according to Claim 1 in which X = N,

 R^{1} , R^{2} , R^{3} = independently of one another CN, OH, COR^{4} or $CH_{2}R^{4}$,

 $R^4 = OH, NH_2, NHB \text{ or } NHB_2,$

Q = CH₂, or CO and, if divalent, also CH,

B = alkyl having 1-6 C atoms,

m = 4 and

n = 0,

and or physiologically acceptable salts, derivatives, solvates and or stereoisomers thereof, including mixtures thereof in all ratios.

- 3. (Currently Amended) A compound Compounds according to Claim 1 which is
 - a) 5-{4-[4-(5-cyano-2,3-dihydro-1H-indol-3-yl)butyl]piperazin-1-yl}benzo-furan-2-carboxamide

b) 5 [4 [4 (5 cyano 6 hydroxy 1H indol-3 yl)butyl]piperazin 1 yl]benzofuran 2 carboxamide.

<u>or</u>

e)b) 5-{4-[4-(5-cyano-2-oxo-2,3-dihydro-1H-indol-3-yl)butyl]piperazin-1-yl}-benzofuran-2-carboxamide.

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- 4. (Currently Amended) Process for the preparation of compounds according to claim 1 and or physiologically acceptable salts, derivatives, solvates and or stereoisomers thereof, characterised in comprising
- a) that reacting a compound of the formula II, in which R¹, R², R³, X, m and n have the meanings indicated in Claim 1,

$$R^{1}$$
 R^{2}
 R^{2}
 R^{3}

is reacted with dimethyl sulfoxide and concentrated HCl, or

b) that reacting a compound of the formula III, in which R¹, R², and n have the meanings indicated in Claim 1, and Y is a halogen, in particular chlorine, or an alcohol provided with protecting groups, known to the person skilled in the art,

is reacted with trifluoroacetic acid and triethylsilane and is subsequently $\frac{1}{1}$ eoupled coupling with a compound of the formula IV, in which R^3 , X and n have the meanings indicated in Claim 1

$$HN$$
 CnH_{2n}
 OR^3
 IV

or

c) that reacting a compound of the formula V, in which R² and m have the meanings indicated in Claim 1 and Y is a halogen, in particular chlorine, or an alcohol provided with protecting groups, known to the person skilled in the art,

$$H_2NOC$$
 R^2
 H_2NOC
 V

is reacted with a dehydrating reagent and is subsequently

- 5. (Canceled)
- 6. (Currently Amended) Pharmaceutical composition comprising at least one compound according to claim 1 and/or physiologically acceptable salts, derivatives, solvates and or stereoisomers thereof, including mixtures thereof in all ratios, and a pharmaceutically acceptable carrier.
- 7. (Original) Pharmaceutical composition, according to Claim 6 comprising further excipients and/or adjuvants.
- 8. (Currently Amended) Pharmaceutical composition comprising at least one compound according to claim 1 and/or physiologically acceptable salts, derivatives, solvates and or stereoisomers thereof, including mixtures thereof in all ratios, and at least one further medicament active ingredient.
- 9. (Currently Amended) Process for the preparation of a pharmaceutical composition, characterised in that comprising bringing a compound according to claim 1 and/or one of its physiologically acceptable salts, derivatives, solvates and or stereoisomers, including mixtures thereof in all ratios, is brought-into a suitable dosage form together with a solid, liquid or semi-liquid excipient or adjuvant.

10. (Canceled)

11. (Currently Amended) Use of compounds according to claim 1 and/or physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, A method for the preparation of a medicament for the treatment of diseases associated with the serotonin receptor and/or serotonin

reuptake, comprising administering to a host in need thereof an effective amount of a compound according to Claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.

- 12. (Currently Amended) Use of compounds according to claim 1 and/or physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament as A method of achieving an anxiolytic, antidepressant, neuroleptic and/or antihypertonic effect and/or for positively influencing obsessive-compulsive disorder (OCD), sleeping disorders, tardive dyskinesia, learning disorders, age-dependent memory disorders, eating disorders, such as bulimia, and/or sexual dysfunctions, comprising administering to a host in need thereof an effective amount of a compound according to claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.
- 13. (Currently Amended) Use of compounds according to claim 1 and/or physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament method for the treatment of psychoses, schizophrenia, schizo-affective psychosis, cyclothymia, epilepsy, cramps, depression (sub-types of severe depression and cyclothymic depression), pathogenic anxiety states (sub-types of panic attacks with or without agoraphobia), superexcitation, hyperactivity, stress illnesses, post-traumatic stress disorders, sleeping disorders, narcolepsy, cyclic manic depression, attention disorders in children and youths, severe developmental disorders, and disorders of social behaviour behavior with mental retardation, obsessive-compulsive disorders in the narrower (OCD) and broader sense (OCSD), addiction disorders, disorders in nutrient uptake or eating disorders, for example bulimia, obesity or anorexia nervosa, fibromyalgia, and psychiatric symptoms in senile dementia and or Alzheimer's-type dementia, cognitive impairments (learning and memory disorders),

in particular age dependent memory disorders, dementia, tardive dyskinesia, neurodegenerative diseases, such as Parkinson's disease, Alzheimer's disease, Huntington's disease, lathyrism, amyotrophic lateral sclerosis, Lewy bodies dementia, Tourette's syndrome, sexual dysfunctions, premenstrual syndrome, acromegaly, hypogonadism, secondary amenorrhoea, undesired puerperal lactation, extrapyramidal motor disorders, for the treatment of side effects arising in the treatment of extrapyramidal motor disorders with conventional anti-Parkinson's medicaments, and of extrapyramidal symptoms (EPS), tension states, side effects of hypertonia treatment induced by neuroleptics (for example with a methyldopa) or for the prophylaxis, treatment and control of cerebral infarctions (apoplexia cerebri), such as strokes and cerebral ischaemia, or for the treatment of pain, in particular chronic pain, migraine, CNS trauma, hypoglycaemia, asthma, glaucoma, cytomegaly and for the treatment of other degenerative retinal diseases, incontinence, tinnitus, or for the treatment of loss of hearing induced by aminoglycoside antibiotics, comprising administering to a host in need thereof an effective amount of a compound according to claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.

14. (Currently Amended) ASet (kit) consisting of separate packs of

- a) an effective amount of a compound according to claim 1 and/or physiologically acceptable salts, derivatives, solvates andor stereoisomers thereof, including mixtures thereof in all ratios, and
- b) an effective amount of a further medicament active ingredient.